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Transmitted herewith for filing is the patent application of

Inventor(s): **Sergey LAMANSKY and Mark E. THOMPSON**

For : **ORGANOMETALLIC PLATINUM COMPLEXES FOR
PHOSPHORESCENCE BASED ORGANIC LIGHT EMITTING
DEVICES**

Enclosed are:

1. 12 sheets of specification, 1 sheet of claims, and 1 sheet of abstract.
2. 8 sheets of drawings.
3. Declaration (unexecuted).
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organometallic platinum complexes
for phosphorescence based
organic light emitting devices

ORGANOMETALLIC PLATINUM COMPLEXES FOR PHOSPHORESCENCE BASED ORGANIC LIGHT EMITTING DEVICES

FIELD OF THE INVENTION

5 The present invention is directed to phosphorescence based organic light emitting devices that have improved electroluminescent characteristics.

BACKGROUND OF THE INVENTION

10 The technology of organic light emitting diodes (OLEDs) is undergoing rapid development. OLEDs originally utilized the electroluminescence produced from electrically excited molecules that emitted light from their singlet states. Such radiative emission from a singlet excited state is referred to as fluorescence. More recent work has demonstrated that higher power efficiency OLEDs can be made using molecules that emit light from their triplet state, defined as phosphorescence.

15 Such electrophosphorescence makes it possible for phosphorescent OLEDs to have substantially higher quantum efficiencies than are possible for OLEDs that only produce fluorescence. This is based on the understanding that the excitons created in an OLED are produced, according to simple statistical arguments as well as experimental measurements, approximately 75% as triplet excitons and 25% as singlet excitons. The triplet excitons more readily transfer their energy to triplet excited states that can produce phosphorescence
20 whereas the singlet excitons typically transfer their energy to singlet excited states that can produce fluorescence. Since the lowest emissive singlet excited state of an organic molecule is typically at a slightly higher energy than the lowest triplet excited state, the singlet excited state may relax, by an intersystem crossing process, to the emissive triplet excited state. This means that all the exciton excitation energy may be converted into triplet state excitation
25 energy, which then becomes available as phosphorescent emission. Thus, electrophosphorescent OLEDs have a theoretical quantum efficiency of 100%, since all the exciton excitation energy can become available as electrophosphorescence.

As a consequence, since the discovery that phosphorescent materials could be used in an OLED, Baldo *et al.*, "Highly Efficient Phosphorescent Emission from Organic Electroluminescent Devices", *Nature*, vol. 395, 151-154, 1998, there is now much interest in finding more efficient electrophosphorescent materials.

Typically phosphorescent emission from organic molecules is less common than fluorescent emission. However, phosphorescence can be observed from organic molecules under an appropriate set of conditions. Organic molecules coordinated to lanthanide elements often phosphoresce from excited states localized on the lanthanide metal. The europium diketonate complexes illustrate one group of these types of species. Organic phosphorescence is also often observed in molecules containing heteroatoms with unshared pairs of electrons at very low temperatures. Benzophenone and 2,2'-bipyridine are such molecules. Phosphorescence can be enhanced over fluorescence is by confining, preferably through bonding, the organic molecule in close proximity to an atom of high atomic number. This phenomenon, called the heavy atom effect, is created by a mechanism known as spin-orbit coupling. A related phosphorescent transition is a metal-to-ligand charge transfer (MLCT) that is observed in molecules such as tris(2-phenylpyridine)iridium(III).

However, molecules that phosphoresce from MLCT states typically emit light that is of lower energy than that observed from the unbound organic ligand. This lowering of emission energy makes it difficult to develop organic molecules that phosphoresce in the technologically useful blue and green colors of the visible spectrum where the unperturbed phosphorescence typically occurs.

It would be desirable if more efficient electrophosphorescent materials could be found, particularly materials that produce their emission in the blue region of the spectrum.

SUMMARY OF THE INVENTION

The present invention is directed to organic light emitting devices, and methods of fabricating the same, that involve using emissive organometallic compounds that produce improved electroluminescence.

Specific embodiments of the present invention are directed to OLEDs using emissive organometallic compounds that produce improved electrophosphorescence in the blue region of the visible spectrum.

The present invention is directed, in addition, to a method of selecting organometallic compounds that have improved electroluminescent properties, for example, in the blue region of the visible spectrum.

The organometallic compounds of the present invention include, in particular, compounds having a metal atom bound to a single carbon-coordination ligand, wherein the carbon-coordination ligand is a mono-anionic carbon-coordination ligand.

A carbon-coordination ligand is a ligand that is bound to the metal atom via a carbon-metal bond. In view of what one skilled in the art might view as a strict definition of organometallic compounds, such as described in Inorganic Chemistry, by Gary L. Miessler and Donald A. Tarr, 2nd edition, Prentice Hall, 1999, the compounds of the present invention are referred to herein as organometallic compounds since these compounds include a metal-carbon bond.

The preferred metals of the present invention are metals that can provide strong spin-orbit coupling of the metal atoms with the carbon-coordination ligand. Such metals include, in particular, the heavy metals having an atomic number of at least 72. Particularly preferred metals include Pt, Ir and Au, with the most preferred being Pt.

The organometallic compounds of the present invention have a single carbon-coordination ligand wherein the single carbon-coordination ligand is a mono-anionic ligand. In particular, the metal atom is bound to only one carbon atom of the carbon-coordination ligand. Thus, while the organometallic compounds that are used in the OLEDs of the present invention may typically include more than one ligand, only one ligand is a carbon-coordination ligand. Thus, the organometallic compounds of the present invention include only one carbon-metal bond.

The carbon-coordination ligand is preferably selected from those ligands that exhibit strong charge transfer absorption characteristics, for example, a molar absorptivity of at least 1,000 L/mole-cm, preferably, at least about 2,000-4,000 L/mole-cm. Such absorption bands involve transfer of electrons from molecular orbitals that are primarily ligand in character to

orbitals that are primarily metal in character or, alternatively, from orbitals that are primarily metal in character to molecular orbitals that are primarily ligand in character. Miessler and Tarr. Such an excitation mechanism results in a charge transfer transition that may be designated as a ligand-to-metal charge transfer (LMCT) or as a metal-to-ligand charge transfer (MLCT), respectively. The former may be characterized as a partial reduction of the metal atom and the latter as a partial oxidation of the metal atom.

Selection of a carbon-coordination ligand to give a high molar absorptivity of the organometallic compound results in an organometallic compound that is capable of providing highly efficient electroluminescence when used in an OLED. However, rather than functioning as strongly absorbing species in the OLED, such organometallic compounds have highly emissive excited states that are produced when a voltage is applied across the OLED. The high molar absorptivities of such ligands may be used to select ligands that produce highly efficient electroluminescence in an OLED. Such ligands may be selected to have empty pi-symmetry orbitals on the ligands that become acceptor orbitals upon absorption of light.

In the preferred embodiments of the present invention, the ligand is selected, in particular, so as to give a strong metal-to-ligand charge transfer (MLCT) absorption band. Such ligands are selected to have empty anti-bonding π^* orbitals on the ligands that become acceptor orbitals upon absorption of light. As representative embodiments of the present invention, the carbon-coordination ligand may be selected from the class of materials such as described, for example, in Comprehensive Coordination Chemistry, Vols. 1-7, G. Wilkinson, Ed., Pergamon Press, 1987.

In addition to being bound to a single mono-anionic carbon-coordination ligand, the metal atom of the organometallic compounds of the present invention is also bound to one or more additional ligands, each of which are all non-carbon-coordination ligands. A non-carbon-coordination ligand is one that does not form any metal-carbon bonds with the metal atom of the organometallic compound. For the preferred embodiments of the present invention for which a metal to ligand charge transfer complex (MLCT) is employed, the non-carbon-coordination ligands are preferably ligands having a strong electrophilic character such that the ligands draw electrons away from the metal atom. Representative non-carbon-coordination ligands may also be selected, for example, from Comprehensive Coordination Chemistry, Vols. 1-7, G. Wilkinson, Ed., Pergamon Press, 1987.

Without intending to be limited to the theory of how the present invention works, it is believed that the improved electroluminescent properties that are observed for the OLEDs of the present invention may be attributed to a combination of factors. For example, it is believed that selection of heavy metals that are capable of forming metal-to-ligand charge transfer (MLCT) states with carbon-coordination ligands that have empty π^* orbitals, such as phosphorescent materials produce highly efficient electrophosphorescent OLEDs. The electroluminescence from representative organometallic compounds of the present invention shows a vibronic fine structure that indicates that the emission is from an excited state that has a wave function represented by a mixture of the MLCT state of the organometallic compound and the excited triplet state of the carbon-coordination ligand. Since the radiative emission is from a triplet excited state, the emission is referred to as phosphorescence.

It is further believed a higher energy radiative emission may be achieved by including electron-withdrawing groups on the carbon-coordination ligand and/or by selecting the non-carbon-coordination ligand to have a strong electron withdrawing character. Without being limited to the precise theory of how the higher energy radiative emissive may be achieved, it is believed that the electron-withdrawing groups tend to remove electron density from the highest occupied molecular orbitals (HOMO) that include the ligand and the metal atom, thus altering the relative energy levels of the ground state and the excited state such that the overall MLCT transition energy from the ground state to the excited state increases. The preferred organometallic compounds of the present invention include, thus, strong electron-withdrawing groups on the carbon-coordination ligand and/or non-carbon-coordination ligands having a strong electron-withdrawing character.

Another aspect of the present invention relates to the discovery that OLEDs incorporating emissive organometallic compounds having a single mono-anionic carbon-coordination ligand have substantially higher external quantum efficiencies than compound with bis-substituted carbon-coordination ligands. For example, for the compound having the chemical structure $[(ppy)Pt(acac)]$ was found to produce strong photophosphorescence at room temperature. In contrast, a compound having the structure $[Pt(ppy)_2]$ was found not to produce any visible photophosphoresce at room temperature.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows the chemical structures of representative organometallic compounds described herein.

Figure 2 shows the emission spectrum of both Pt(ppy)₂ and Pt(ppy)₂Br₂. The former gives green emission, partly from MLCT transitions, and the latter gives blue emission, predominantly from a triplet π - π^* transition. The structure observed for the Pt(ppy)₂Br₂ spectrum is consistent with ligand-centered emission. The luminescent lifetimes for the two complexes are 4 and 150 microseconds.

Figure 3 is a plot showing the emission spectra of (ppy)AuCl₂ and (ppy)Au(2,2'-biphenylene). Both emit from ligand triplet π - π^* transitions.

Figure 4 is a CIE diagram providing the coordinates of (C-N)Pt(acac) complexes. All coordinates are based on solution photoluminescent measurements except for 4,5-F₂ppy-EL, which corresponds to the electroluminescent spectrum. The Ir(ppy)₃ is an electroluminescent spectrum as well.

Figure 5 is a plot depicting the photoluminescent emission spectra of (4,6-F₂ppy)Pt(acac) at room temperature (RT) and at 77 K. Also shown are the excitation spectra taken at 77 K and the absorbance spectra taken at room temperature for the same complex.

Figure 6 illustrates the normalized photoluminescent emission spectra of (ppy)Pt(acac), (4,5 dfppy)Pt(acac), and (4,5 dfppy)Pt(pico).

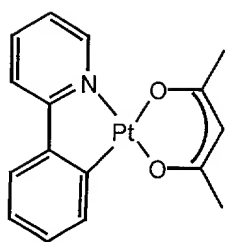
Figure 7 illustrates the normalized photoluminescent emission spectra of typPy(acac), bzqPt(acac), and btpPt(acac).

Figure 8 illustrates the normalized electroluminescent emission spectra for OLEDs prepared with (2-(4,5-F₂phenyl)pyridinato)platinum(acetyl acetate). The OLEDs had a ITO/PVK-PBD-dopant/Alq₃/Mg-Ag layer structure. The PVK layer was deposited as a single, homogeneous layer by spin coating. PVK = polyvinylcarbazole and PBD = (4-biphenyl)(4-tertbutyl)oxiadiazole. The Alq₃ and Mg-Ag layers were deposited by thermal evaporation. The OLED had an external efficiency of 1.3 % and a turn on voltage of 5 Volts. The spectra of the EL output and as well as the PL signal are shown.

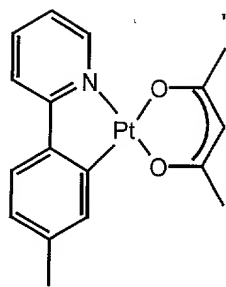
DETAILED DESCRIPTION OF THE INVENTION

The present disclosure is aimed at a new class of platinum complexes, which give efficient phosphorescence. The representative complexes have a single organometallic ligand (cyclometallated) and a bidentate coordination ligand (such as acetylacetonate). Several examples of these complexes are given below along with their spectra. The emission spectra from these complexes show vibronic fine structure, consistent with strong ligand $\pi - \pi^*$ character in the phosphorescent transition. Strong ligand $\pi - \pi^*$ character is also consistent with the fact that the emission energy is strongly dependent on the identity of the ligand, as shown in the spectra shown below. Emission from these complexes results from a mixture of metal to ligand charge transfer (MLCT) and ligand based transitions. The MLCT is critical to enhance the efficiency of intersystem crossing and phosphorescence. The emission is dominated by the cyclometallated ligand and the MLCT between the Pt ion and that ligand. The emission spectrum is only slightly affected by changing the acetylacetonate ligand (acac) to a picolinic acid (pico), as shown for the (ppy)PtX complexes below. This minor shift most likely occurs due to a shift in the Pt based HOMO level by the pico ligand, leading to a red shift in the MLCT and a corresponding red shift in the emission spectrum.

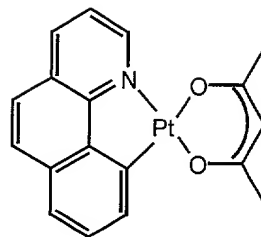
One of these complexes, i.e. (2-(4,5-F₂-phenyl)pyridinato)Pt(acetylacetonate), was used as a phosphorescent dopant in a polymer OLED and gave an emission spectrum identical to the photoluminescence spectrum and an external quantum efficiency of 1.3%.



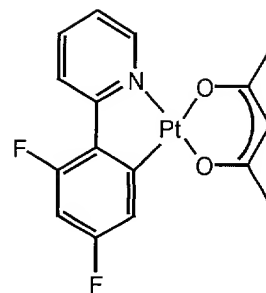
(ppy)Pt(acac)



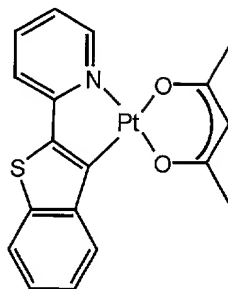
(tpy)Pt(acac)



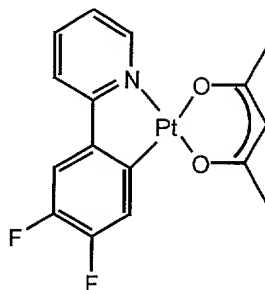
(bzq)Pt(acac)



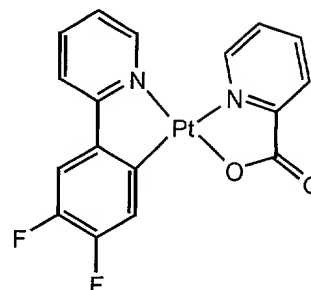
(4,6-F₂ppy)Pt(acac)



(btp)Pt(acac)



(4,5-F₂ppy)Pt(acac)



(4,5-F₂ppy)Pt(pico)

As representative electrophosphorescent compounds of the present invention, the carbon-coordination ligand forms a cyclometallated ring that includes the organometallic carbon-metal bond and a dative bond between the metal atom and a nitrogen, sulfur or oxygen group, for example, Pt(II)-(2-phenylpyridinato-N,C^{2'})(acetyl acetate), herein referred to as Pt(ppy)(acac) or (ppy)Pt(acac). The carbon atom that is bound to the metal may be present as part of a substituted or unsubstituted, saturated hydrocarbon; a substituted or unsubstituted, aromatic system, for example, phenylene or naphthalene compounds; or a substituted or unsubstituted heterocyclic system, which might include, for example, substituted or unsubstituted thiophenes, furans, pyridines and pyrroles. The group in the cyclometallated ring that forms a dative bond with the metal atom may be independently selected also to include a substituted or unsubstituted, saturated hydrocarbon; a substituted or unsubstituted, aromatic system, for example, phenylene or naphthalene compounds; or a substituted or unsubstituted heterocyclic system, which might include, for example, thiophenes, furans, pyridines and pyrroles.

EXAMPLES OF THE INVENTION

All procedures involving K_2PtCl_4 or any other Pt species were carried out in inert gas atmosphere in spite of the air stability of the compounds, the main concern being their oxidative stability and stability of intermediate complexes at high temperatures used in the reactions. NMR spectra were recorded on Bruker AMX 360 MHz or 500 MHz instruments unless specified otherwise. Solid probe MS spectra were taken with Hewlett Packard GC/MS instrument with electron impact ionization and model 5873 mass sensitive detector. High resolution mass spectrometry was done at Frik Chem Laboratories at Princeton University. Elemental analysis data was recorded at the Microanalysis Laboratory at the University of Illinois, Urbana-Champaign.

Pt(II) μ -chlorobridged dimers of the structure $[Pt(C-N)(\mu-Cl)_2Pt(C-N)]$ containing cyclometalated carbon,nitrogen ligands (C,N) used in the study were prepared according to Cave G.W.V., Fanizzi F.P., Deeth R.J., Errington W., Rourke J.P., Organometallics 2000, 19, 1355.

Platinum(II) (2-phenylpyridinato- N,C^2') (acetyl acetate) $[Pt(ppy)(acac)]$. 100 mg of $Pt(ppy)(\mu-Cl)_2Pt(ppy)$ dimer, 25 mg of 2,4-pentanedione and 85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright yellow-green solid (36% yield). 1H NMR (360 MHz, acetone- d_6), ppm: 9.00 (d, 1H, J 5.8Hz), 8.02 (dt, 1H, J 1.6, 7.4Hz), 7.89 (d, 1H, J 7.9Hz), 7.57 (dd, 1H, J 1.6, 7.4Hz), 7.51 (dd, 1H, J 1.6, 7.9Hz), 7.32 (dt, 1H, J 1.6, 6.8Hz), 7.11 (dt, 1H, J 1.6, 7.9Hz), 7.04 (dt, 1H, J 1.6, 7.4Hz), 5.55 (s, 1H), 1.96 (s, 3H), 1.95 (s, 3H). See Figure 4, compound number 5. See also Figure 1(a).

Platinum(II) (2-(p-tolyl)pyridinato- N,C^2') (acetyl acetate) $[Pt(tpy)(acac)]$. 100 mg of $Pt(tpy)(\mu-Cl)_2Pt(tpy)$ dimer, 25 mg of 2,4-pentanedione and 85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright yellow-green solid (42% yield). 1H NMR (360 MHz, $CDCl_3$), ppm: 8.94 (d, 1H, J 5.9Hz), 7.74 (t, 1H, J 6.8Hz), 7.53 (d, 1H, J 7.8Hz), 7.39 (s, 1H), 7.30 (d, 1H, J 7.8Hz), 7.04 (t, 1H, J

6.8Hz), 6.88 (d, 1H, J 7.8Hz), 5.45 (s, 1H), 2.00 (s, 3H), 1.98 (s, 3H), 1.95 (s, 3H). See Figure 4, compound number 1. See also Figure 1(b).

5 *Platinum(II) (7,8-benzoquinolino-*N,C*^{3'}) (acetyl acetonate) [Pt(bzq)(acac)]*. 100 mg of Pt(bzq)(μ-Cl)₂Pt(bzq) dimer, 25 mg of 2,4-pentanedione and 85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright yellow solid (27% yield). ¹H NMR (360 MHz, acetone-d₆), ppm: 9.13 (d, 1H, J 5.4Hz), 8.25 (d, 1H, J 8.3Hz), 7.75 (m, 2H), 7.50-7.57 (m, 3H), 7.44 (dd, 1H, J 5.4, 5.4Hz), 5.52 (s, 1H), 10 2.04 (s, 6H). See Figure 4, compound number 2. See also Figure 1(c).

15 *Platinum(II) (2-benzylpyrino-*N,C*^{2'}) (acetyl acetonate) [Pt(bzpy)(acac)]*. 100 mg of Pt(bzpy)(μ-Cl)₂Pt(bzpy) dimer, 25 mg of 2,4-pentanedione and 85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright yellowish green solid (20% yield). ¹H NMR (500 MHz, CDCl₃), ppm: 8.88 (d, 1H), 7.71 (t, 1H), 7.35-7.43 (m, 2H), 7.13 (t, 1H), 6.98-7.02 (m, 2H), 6.91 (t, 1H), 5.49 (s, 1H), 4.16 (s, 2H), 1.96 (s, 3H), 1.95 (s, 3H).

20 *Platinum(II) (2-(2'-thienyl)pyridino-*N,C*^{3'}) (acetyl acetonate) [Pt(thpy)(acac)]*. 100 mg of Pt(thpy)(μ-Cl)₂Pt(thpy) dimer, 25 mg of 2,4-pentanedione and 85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright orange solid (20% yield). ¹H NMR (500 MHz, CDCl₃), ppm: 8.78 (d, 1H), 7.67 (t, 1H), 7.46 (d, 1H), 7.26 (d, 1H), 7.17 (d, 1H), 6.86 (t, 1H), 5.46 (s, 1H), 1.98 (s, 3H), 1.95 (s, 3H). 25 (s, 3H).

30 *Platinum(II) (2-(2'-(4',5'-benzothienyl)pyridino-*N,C*^{3'}) (acetyl acetonate) [Pt(btp)(acac)]*. 100 mg of Pt(btp)(μ-Cl)₂Pt(btp) dimer, 25 mg of 2,4-pentanedione and 85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography

(silica/dichloromethane) gave orange-red solid (20% yield). ^1H NMR (360 MHz, CDCl_3), ppm: 8.90 (d, 1H, J 5.9Hz), 8.75-8.79 (m, 1H), 7.77-7.81 (m, 1H), 7.71 (dt, 1H, J 1.5, 7.8Hz), 7.27-7.34 (m, 3H), 6.95 (dt, 1H, J 1.5, 6.8Hz), 5.54 (s, 1H), 2.08 (s, 3H), 2.01 (s, 3H). See Figure 4, compound number 3. See also Figure 1(e).

5 *Platinum(II) (2-(4',6'-difluorophenyl)pyridinato- N,C^2') (acetyl acetonate) [Pt(4,6- F_2 ppy)(acac)].* 131 mg of Pt(4,6- F_2 ppy)($\mu\text{-Cl}$) $_2$ Pt(4,6- F_2 ppy) dimer, 43 mg of 2,4-pentanedione and 109 mg of anhydrous sodium carbonate were refluxed at 100 °C in 10 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash
10 chromatography (silica/dichloromethane) gave bright yellow solid. ^1H NMR (360 MHz, acetone- d_6), ppm: 9.06 (dt, 1H, J 1.0, 5.9Hz), 8.08-8.13 (m, 1H), 8.01 (dt, 1H, J 1.5, 8.3Hz), 7.38-7.43 (m, 1H), 7.05 (dd, 1H, J 2.4, 9.3Hz), 6.69-6.76 (m, 1H), 5.61 (s, 1H), 2.01 (s, 3H), 1.99 (s, 3H). See Figure 4, compound number 4. See also Figure 1(d).

15 *Platinum(II) (2-(4',5'-difluorophenyl)pyridinato- N,C^2') (acetyl acetonate) [Pt(4,5- F_2 ppy)(acac)].* 68 mg of Pt(4,5- F_2 ppy)($\mu\text{-Cl}$) $_2$ Pt(4,5- F_2 ppy) dimer, 36 mg of 2-picolinoc acid and 57 mg of anhydrous sodium carbonate were refluxed at 100 °C in 5 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash
20 chromatography (silica/dichloromethane) gave bright yellow solid. ^1H NMR (360 MHz, acetone- d_6), ppm: 8.99 (d, 1H, J 5.7Hz), 8.06 (dt, 1H, J 2.3, 8.0Hz), 7.90 (d, 1H, J 8.0Hz), 7.62-7.68 (m, 1H), 7.37 (tt, 1H, J 1.7, 5.7Hz), 7.20-7.25 (m, 1H), 5.58 (s, 1H), 1.99 (s, 3H), 1.98 (s, 3H). See Figure 4, compound number 6. See also Figure 1(f).

25 *Platinum(II) (2-(4',5'-difluorophenyl)pyridinato- N,C^2') (2-picolinato) [Pt(4,5- F_2 ppy)(pico)].* 69 mg of Pt(4,5- F_2 ppy)($\mu\text{-Cl}$) $_2$ Pt(4,5- F_2 ppy) dimer, 30 mg of 2-picolinoc acid and 52 mg of anhydrous sodium carbonate were refluxed at 100 °C in 5 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash
30 chromatography (silica/dichloromethane) gave bright yellow solid. ^1H NMR (500 MHz, CDCl_3), ppm: 9.15 (d, 1H, J 5.6Hz), 9.05 (d, 1H, J 5.6Hz), 8.08-8.21 (m, 2H), 7.89 (td, 1H, J 1.2, 8.0Hz), 7.68-7.71 (m, 1H), 7.54 (d, 1H, J 8.0Hz), 7.32-7.36 (m, 1H), 7.12-7.20 (m, 2H). See Figure 4, compound number 7. See also Figure 1(g).

Platinum(II) (2-(4'-cyanophenyl)pyridinato- $N,C^{2'}$) (acetyl acetate)
[Pt(cppy)(acac)]. 69 mg of Pt(cppy)(μ -Cl)₂Pt(cfppy) dimer, 58 mg of 2-picolinoc acid and 52
mg of anhydrous sodium carbonate were refluxed at 100 °C in 5 ml of 2-ethoxyethanol under
inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and
filtration yielded crude product that after drying and flash chromatography
(silica/dichloromethane) gave bright yellow solid. ¹H NMR (360 MHz, acetone- d_6), ppm:
9.07 (dt, 1H, J 1.0, 5.9Hz), 8.14 (dt, 1H, J 1.5, 7.8Hz), 8.05 (dt, 1H, J 1.0, 8.3Hz), 7.77-7.79
(m, 2H), 7.46-7.50 (m, 1H), 7.43 (dd, 1H, J 1.5, 8.3Hz), 5.61 (s, 1H), 2.01 (s, 6H).

OLED preparation and testing. Polymer blend OLEDs were spun coat from
chloroform solution on patterned pre-cleaned and oxygen plasma treated indium tin oxide
(ITO) coated glass substrates and covered with vacuum-deposited aluminum(III) tris(8-
hydroxyquinolate) and/or Mg:Ag (10:1 weight ratio) cathode (500 Å) for the single-layer
and heterostructure architectures, respectively. Typically, 7.5 ml of a chloroform solution
contained 100 mg of PVK, 40 mg of PBD and 2.5 mg of (45F₂ppy)Pt(acac). Chosen spin-
coating conditions (3000 RPM, 40 s, Specialty Coating Systems, Inc.) led to 1300±20 Å-thick
PVK:PBD:dye films as determined by ellipsometry (Rudolph automatic ellipsometer
equipped with a He:Ne laser). Prior to spinning, the solutions were filtered through a 0.2 µm
filter. Tris(8-hydroxyquinoline) aluminum (III) (Sigma-Aldrich, Inc) (Alq₃) was sublimed
prior to use. All measurements on the devices were carried out in air at room temperature.
Device current-voltage and light intensity characteristics were measured using the
LabVIEW™ program by National Instruments with a Keithley 2400 SourceMeter/2000
Multimeter coupled to a Newport 1835-C Optical Meter. Electroluminescence spectra were
recorded at room temperature on a PTI QuantaMaster™ Model C-60SE spectrofluorometer.

Other methods known to those skilled in the art of fabricating OLEDs may be used.

While the invention has been described in detail with reference to certain
embodiments, it will be recognized by those skilled in the art that there are other
embodiments of the invention within the spirit and scope of the claims.

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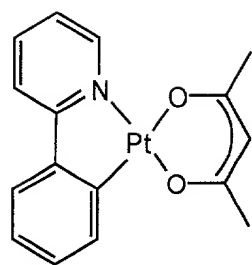
5

- 3.

ABSTRACT

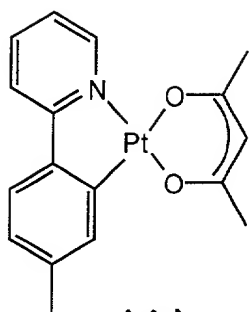
A device for producing electroluminescence comprising an organic light emitting device including an emissive layer comprising an organometallic compound comprised of a metal bound to a single carbon-coordination ligand, with the single carbon-coordination

5 ligand being a monanionic carbon-coordination ligand.



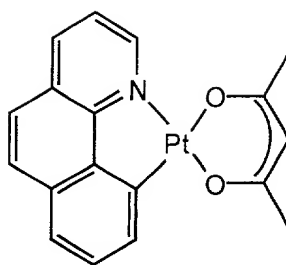
1(a)

(ppy)Pt(acac)



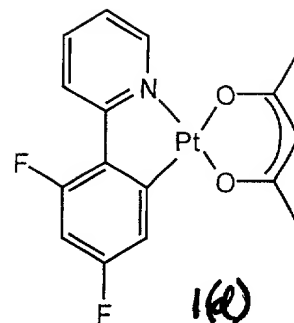
1(b)

(tpy)Pt(acac)



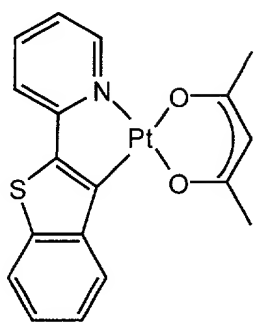
1(c)

(bzq)Pt(acac)



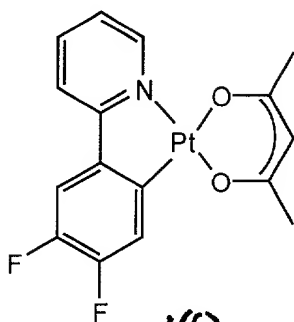
1(d)

(4,6-F₂ppy)Pt(acac)



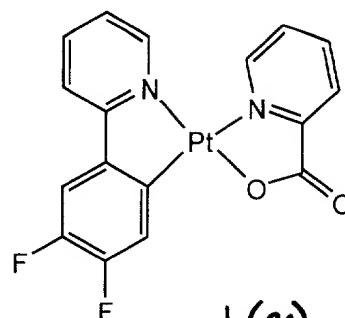
1(e)

(btp)Pt(acac)



1(f)

(4,5-F₂ppy)Pt(acac)



1(g)

(4,5-F₂ppy)Pt(pico)

Figures 1(a) - 1(g)

Figure 2: This Emission spectrum shows the spectra of both Pt(ppy)_2 and $\text{Pt(ppy)}_2\text{Br}_2$. The former gives green emission, partly from MLCT transitions, and the latter gives blue emission, predominantly from a triplet $\pi\text{-}\pi^*$ transition. The structure observed for the $\text{Pt(ppy)}_2\text{Br}_2$ spectrum is consistent with ligand centered emission. The luminescent lifetimes for the two complexes are 4 and 150 μsec .

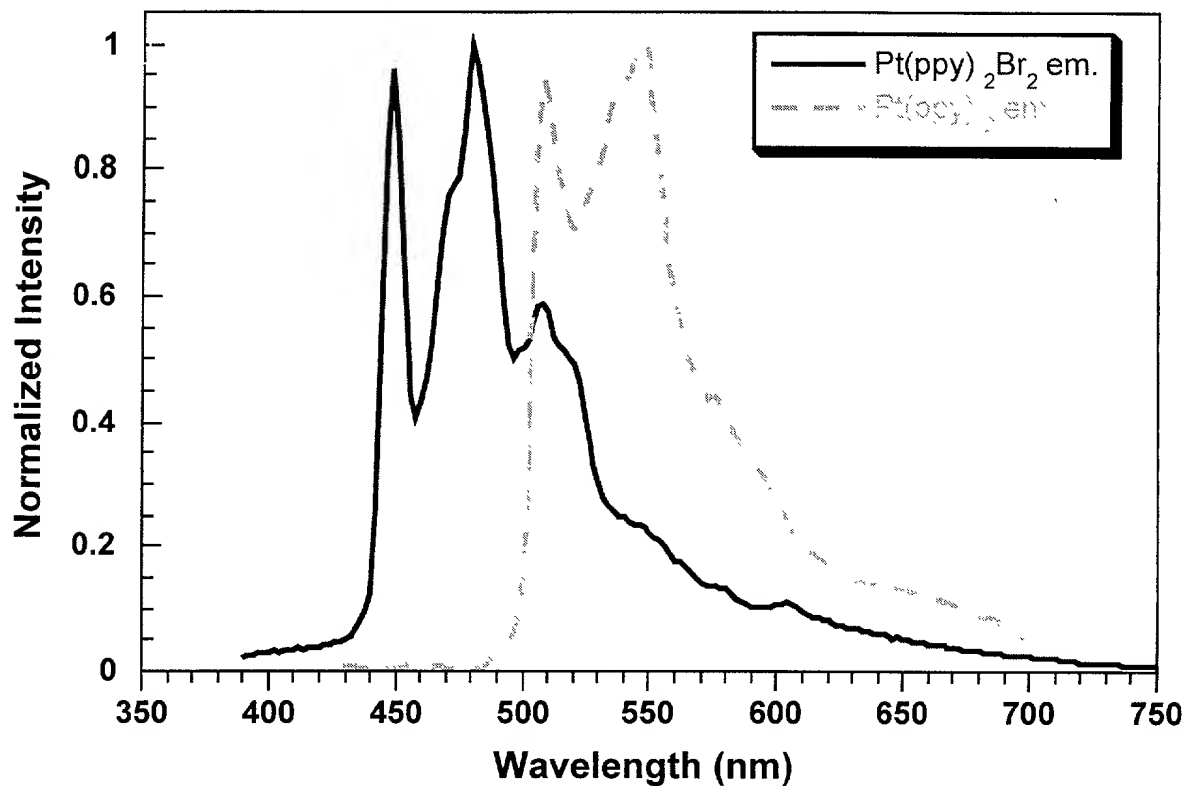


Figure 2

Figure 3: This plot shows the emission spectra of (ppy)AuCl₂ and (ppy)A₂ (2,2'-biphenylene). Both emit from ligand triplet π - π^* transitions.

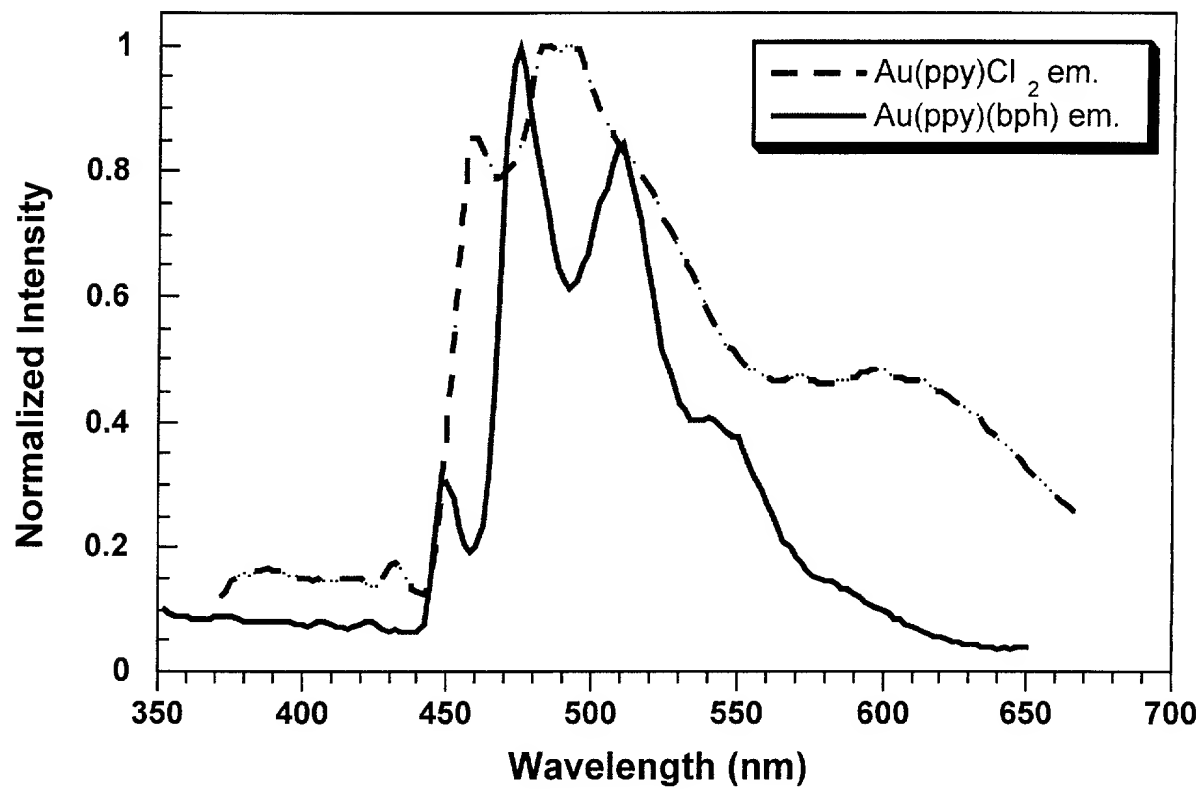


Figure 3

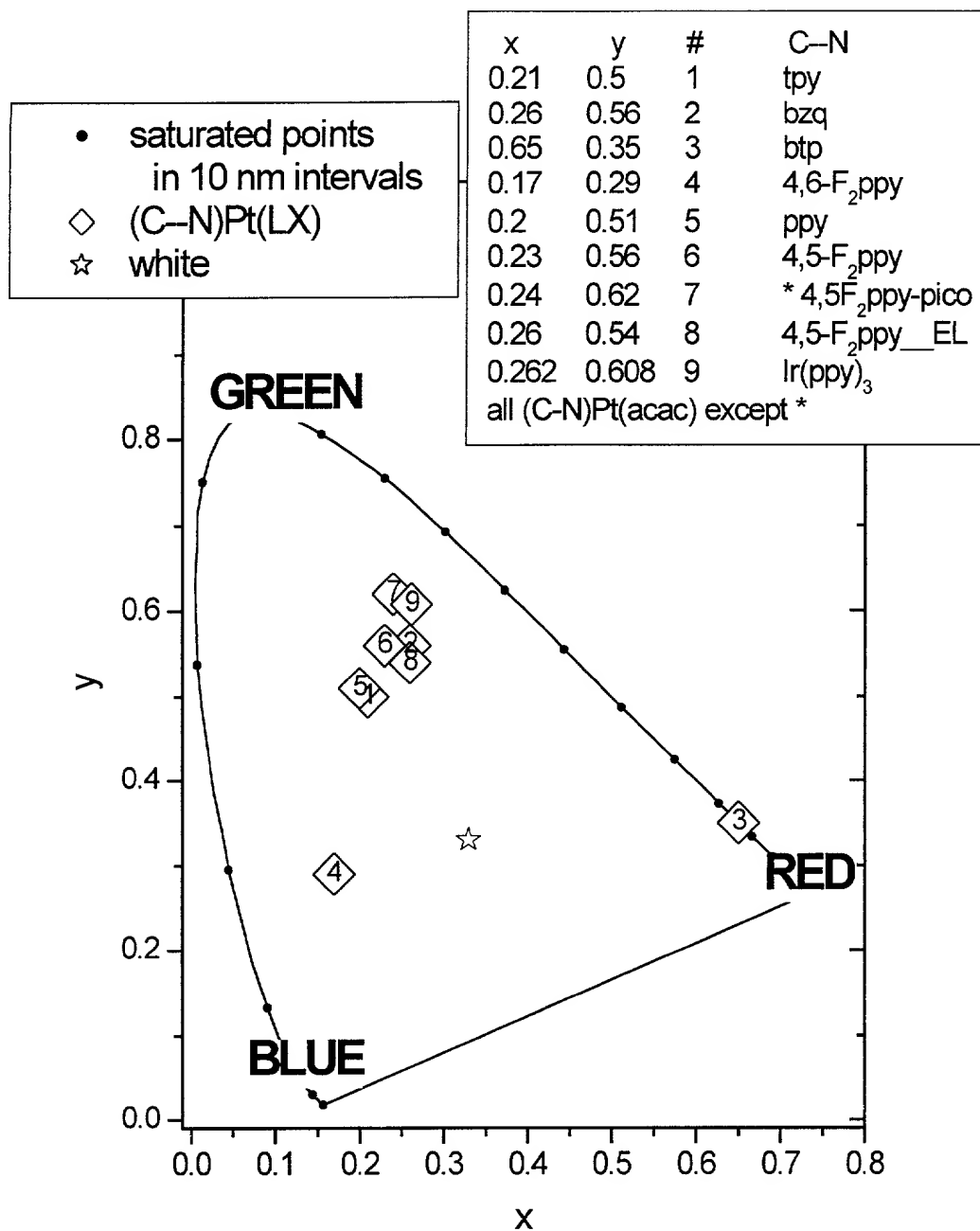


Figure 4

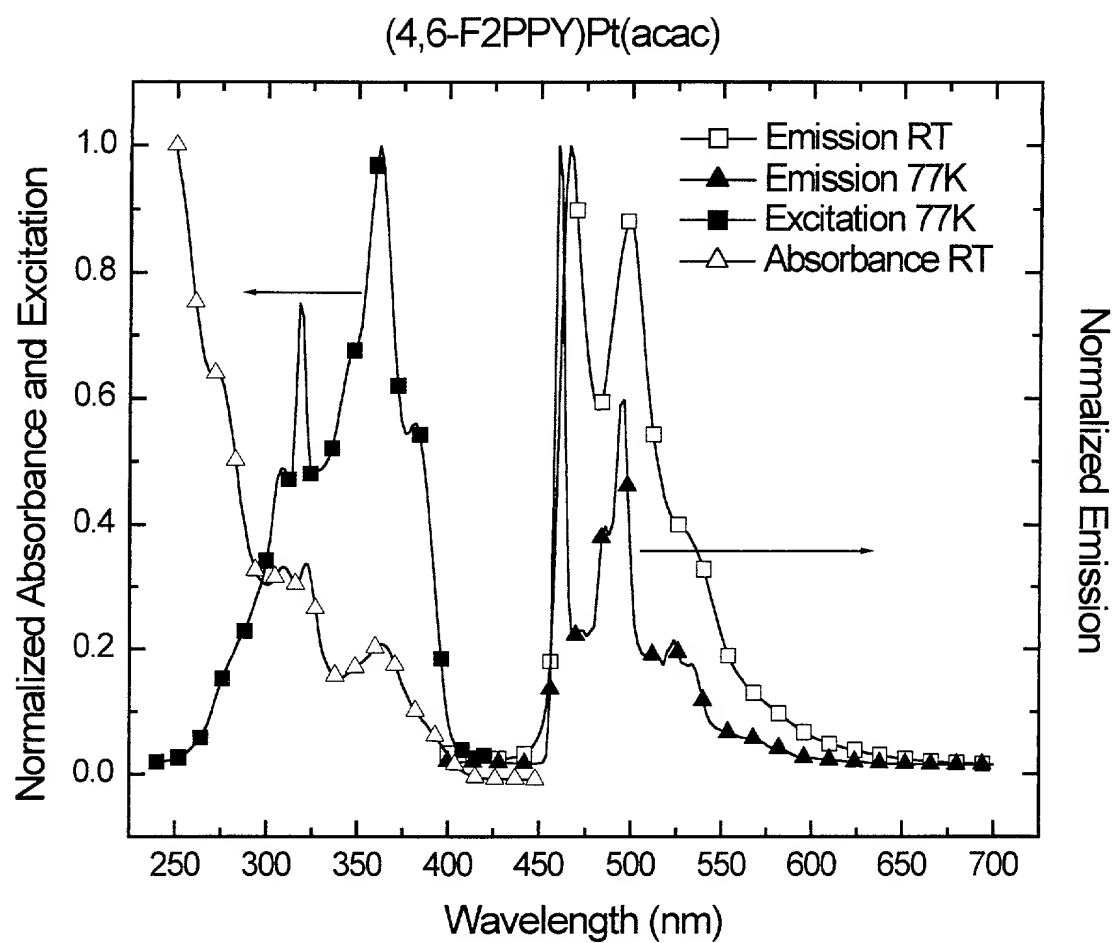


Figure 5

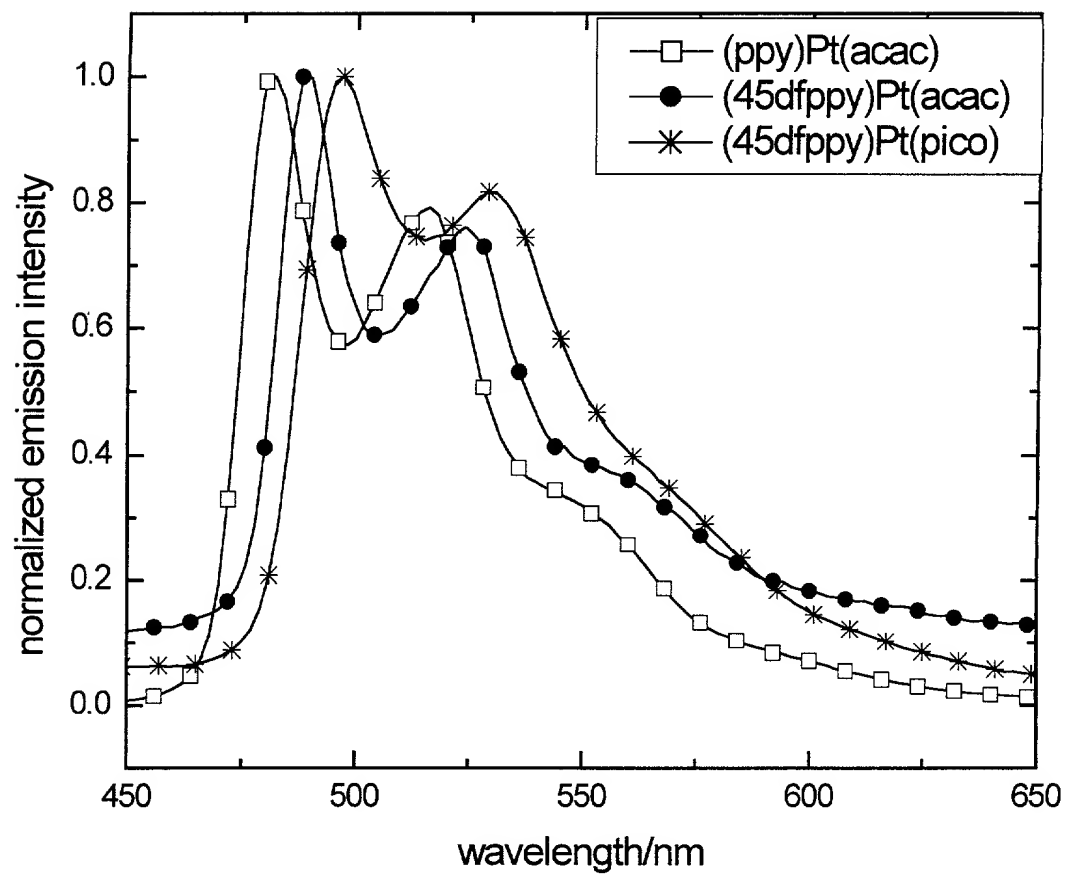


Figure 6

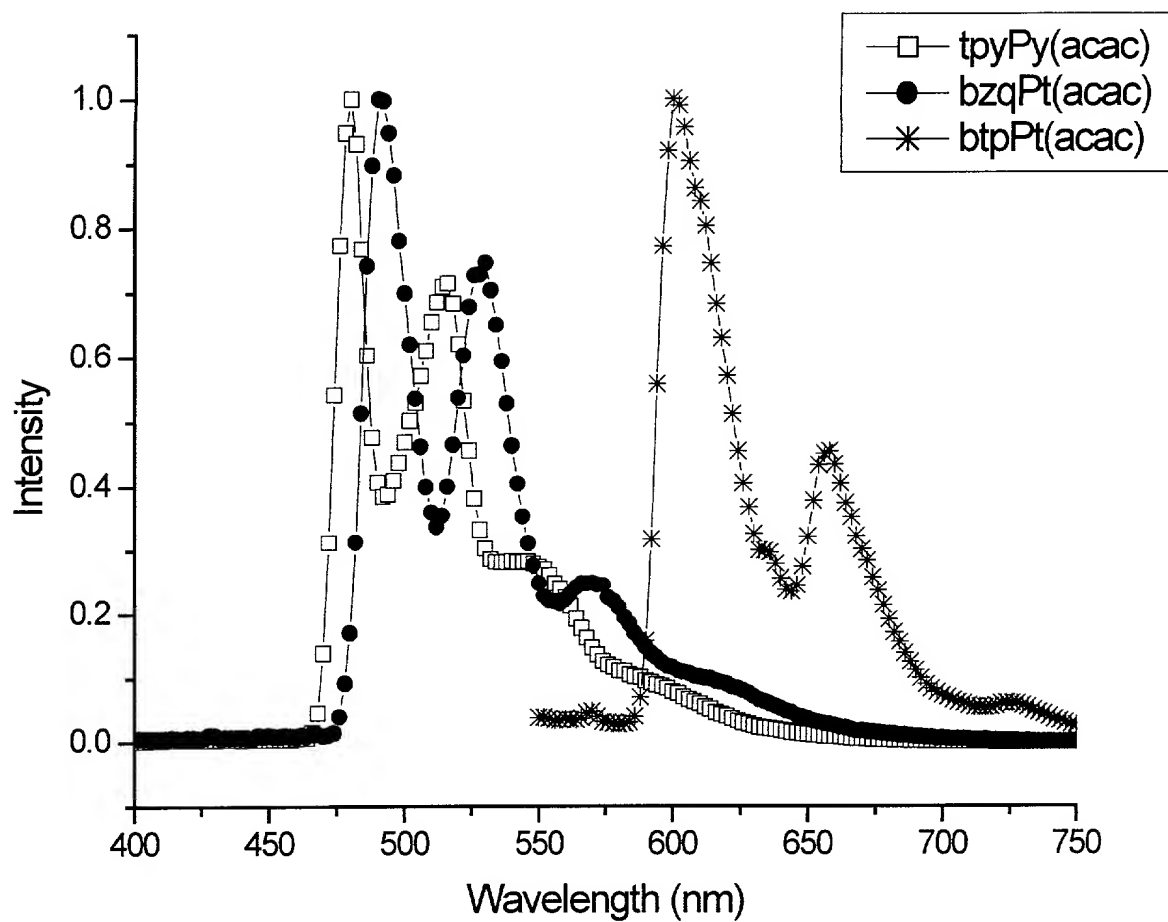


Figure 7

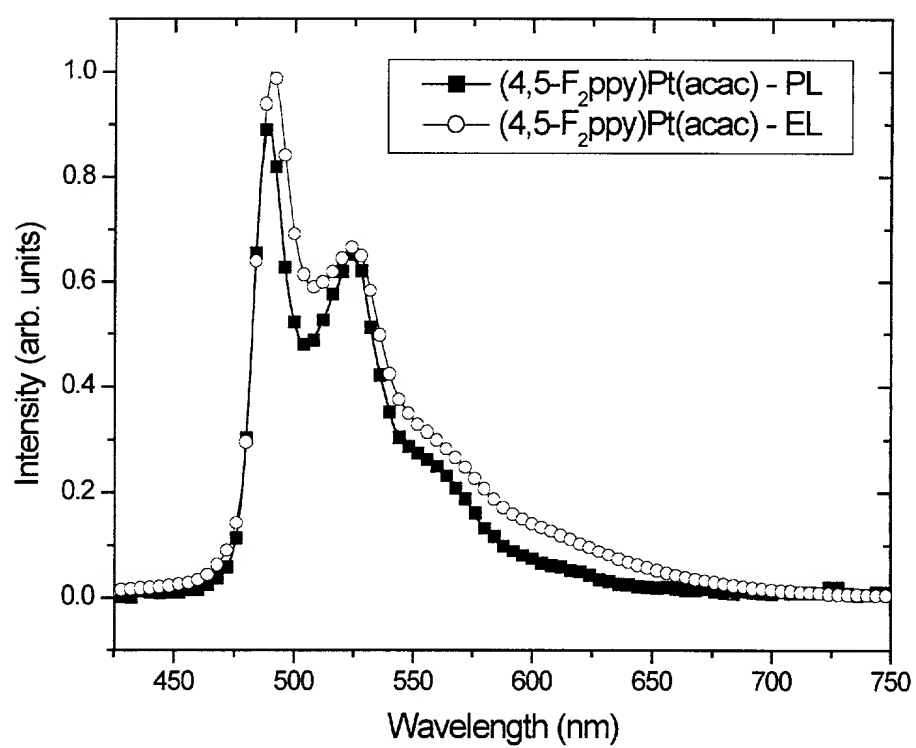


Figure 8

U.S. DEPARTMENT OF COMMERCE
PATENT AND TRADEMARK OFFICE

DECLARATION

ATTORNEY'S DOCKET NO.
10020/20701

As a below named inventor, I hereby declare that:

My residence, post office address, and citizenship are as stated below next to my name,

I believe I am an original, first, and joint inventor of the subject matter that is claimed and for which a patent is sought on the invention entitled **ORGANOMETALLIC PLATINUM COMPLEXES FOR PHOSPHORESCENCE BASED ORGANIC LIGHT EMITTING DEVICES**, the specification of which is being filed herewith.

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, § 1.56(a).

SEND CORRESPONDENCE, AND DIRECT TELEPHONE CALLS TO:

THOMAS F. MEAGHER

KENYON & KENYON
One Broadway
New York, New York 10004
(212) 425-7200 (phone)
(212) 425-5288 (facsimile)

I declare that all statements made herein of my own knowledge are true and all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under § 1001 of Title 18 of the United States Code and that such willful statements may jeopardize the validity of the application or any patent issuing thereon.

EL 32754876505

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Signature		Date	

FULL NAME OF INVENTOR	FAMILY NAME THOMPSON	FIRST GIVEN NAME Mark	SECOND GIVEN NAME E.
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Signature		Date	